

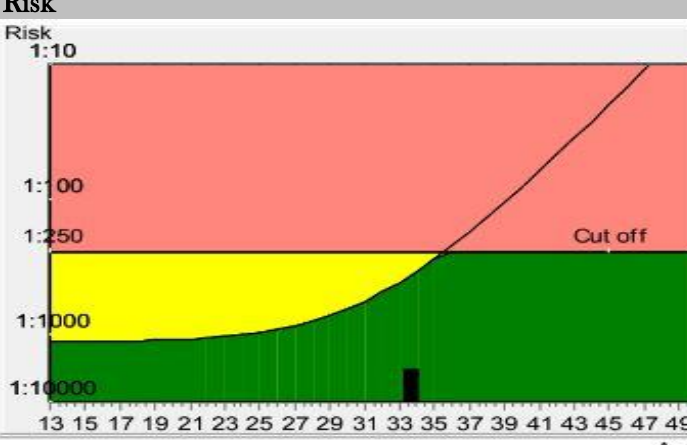
Date of Report 27/12/19
PRISCA 5.0.2.37

Patient Data	
Name	Mrs Anita
Birthdate	2/5/1986
Age at delivery	33.7
Gestational age	13+0
Patient ID	011912250275
Sample ID	10528936
Sample Date	25/12/19

Correction factors	
Fetuses	1 IVF
Weight in kg	45 Diabetes
Smoker	no Origin
unknown	Previous trisomy 21
no	Pregnancies
Asian	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+6
PAPP-A	5.61 mIU/ml	0.76	Method	CRL (<>Robinson)
fb-hCG	78.07 ng/ml	1.59	Scan Date	25/12/19

Risks at sampling date		Down's Syndrome Risk (Trisomy 21 Screening)	
Age Risk	1:366	Crown Rump Length (mm)	66.2
Biochemical Trisomy 21 Risk	1:410	Nuchal translucency MoM	0.56
Combined Trisomy 21 Risk	1:2341	Nasal Bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR PAULDEEP KAUR
		Qualification in measuring NT	MBBS, MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
	<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2341 women with the same data, there is one woman with a trisomy 21 pregnancy and 2340 women with not affected pregnancies.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical approaches and have no diagnostic value!</p>

Trisomy 13/18 + NT	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values
The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.	

